

(11) EP 1 160 348 A2

(12)

EUROPEAN PATENT APPLICATION

(43) Date of publication: 05.12.2001 Bulletin 2001/49

(51) Int CI.7: C23C 4/02

(21) Application number: 01112406.2

(22) Date of filing: 21.05.2001

(84) Designated Contracting States:

AT BE CH CY DE DK ES FI FR GB GR IE IT LI LU

MC NL PT SE TR

Designated Extension States:

AL LT LV MK RO SI

(30) Priority: 22.05.2000 US 576439

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(54) Process for producing graded coated articles

(57) The invention relates to a novel thermal spray process for the deposition of coatings with a graded or layered composition and the coated articles produced thereby. More particularly, the invention relates to feeding mixtures of coating materials to a thermal spray de-

vice and continuously or intermittently changing the composition of the deposited coatings by changing the thermal spray operating parameters. The continuous or Intermittent change in the composition of the coating material during deposition creates a graded or layered coating structure.

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Description

Technical Field

[0001] The invention relates to a novel thermal spray process for the deposition of coatings with a graded or layered composition on a substrate and the coated articles produced thereby. More particularly, the invention relates to feeding at least two coating materials to a thermal spray device and continuously or intermittently changing the composition of the deposited coatings by changing the thermal spray operating parameters. The change in the composition of the coating during deposition creates a graded or layered coating structure.

Background Art

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[0002] The family of thermal spray processes includes detonation gun deposition, high velocity oxy-fuel deposition and its variants such as high velocity air-fuel, plasma spray, flame spray, and electric wire arc spray. In most thermal coating processes metallic, ceramic, cermet, or some polymeric materials in powder, wire or rod form is heated to near or somewhat above its melting point and droplets of the material accelerated in a gas stream. The droplets are directed against the surface of a substrate to be coated where they adhere and flow into thin lamellar particles called splats. The coating is built up of multiple splats overlapping and interlocking. These processes and the coatings they produce have been described in detail in the following: "Advanced Thermal Spray Deposition Techniques", R.C. Tucker, Jr., in Handbook of Deposition Technologies for Films and Coatings, R.F. Bunshah, ed., Second Edition, Noyes Publications, Park Ridge, New Jersey, 1994, pp. 591 to 6421; "Thermal Spray Coatings", R.C. Tucker, Jr. in Handbook of Thin Films Process Technology, Institute of Physics Publishing, Ltd., London, 1995; and "Thermal Spray Coatings", R.C. Tucker, Jr., in Surface Engineering ASM Handbook, Vol. 5, ASM International, Materials Park, Ohio, 1994, pp 497-509.

[0003] In virtually all of the thermal spray processes two of the most important parameters controlling the structure and properties of the coatings are the temperature and velocity of the individual particles as they impact on the surface to be coated. Of these, the temperature of the particles is of greatest importance relative to the present invention. The temperature the particles achieve during the deposition process is a function of a number of parameters including the temperature and enthalpy (heat content) of the process gases, the specific mechanisms of heat transfer to the particles, the composition and thermal properties of the particles, the size and shape distributions of the particles, the mass flow rate of the particles relative to the gas flow rate, and the time of transit of the particles. The velocity the particles achieve is a function of a number of parameters as well, and some of these are the same as those that affect the particle temperature including the composition, velocity and flow rate of the gases, the size and shape distributions of the particles, the mass injection rate and density of the particles. Thus the thermal gas dynamics characteristics of the thermal spray process determine the quality of the resulting coating.

[0004] In a typical detonation gun deposition process, a mixture of oxygen and a fuel such as acetylene along with a pulse of powder of the coating material is injected into a barrel, such as a barrel of about 25 mm in diameter and over a meter long. The gas mixture is detonated, and the detonation wave moving down the barrel heats the powder to near or somewhat above its melting point and accelerates it to a velocity of about 750 m/sec. The molten, or nearly molten droplets of material strike the surface of the substrate to be coated and flow into strongly bonded splats. After each detonation, the barrel is generally purged with an inert gas such as nitrogen, and the process repeated many times a second. Detonation gun coatings typically have a porosity of less than two volume percent with very high cohesive strength as well as very high bond strength to the substrate. In the Super D-Gun™ coating process, the gas mixture includes other fuel gases in addition to acetylene. As a result there is an increase in the volume of the detonation gas products which increases the pressure and hence greatly increases the gas velocity. This, in turn, increases the coating material particle velocity which may exceed 1000 m/sec. The increased particle velocity can result in an increase in coating bond strength, density, and an increase in coating compressive residual stress. In both the detonation gun and Super D-Gun coating processes, nitrogen or another inert gas can be added to the detonation gas mixture to control the temperature of the detonated gas mixture and hence the powder temperature. A number of parameters can be used to control both the particle temperature and velocity including the composition and flow rates of the gases into the gun.

[0005] In high velocity oxy-fuel and related coating processes, an oxygen, air or another source of oxygen is used to burn a fuel such as hydrogen, propane, propylene, acetylene or kerosene in a combustion chamber and the gaseous combustion products allowed to expand through a nozzle. The gas velocity may be supersonic. Powdered coating material is injected into the nozzle and heated to near or above its melting point and accelerated to a relatively high velocity, such as up to about 600 m/sec. for some coating systems. The temperature and velocity of the gas stream through the nozzle, and ultimately the powder particles, can be controlled by varying the composition and flow rate of the gases or liquids into the gun. The molten particles impinge on the surface to be coated and flow into fairly densely packed splats that are well bonded to the substrate and each other.

[0006] In the plasma spray coating process a gas is partially ionized by an electric arc as it flows around a tungsten cathode and through a relatively short converging and diverging nozzle. The partially ionized gas, or gas plasma, is usually based on argon, but may contain, for example, hydrogen, nitrogen, or helium. The temperature of the plasma at its core may exceed 30,000 K and the velocity of the gas may be supersonic. Coating material, usually in the form of powder, is injected into the gas plasma and is heated to near or above its melting point and accelerated to a velocity that may reach about 600 m/sec. The rate of heat transfer to the coating material and the ultimate temperature of the coating material are a function of the flow rate and composition of the gas plasma as well as the torch design and powder injection technique. The molten particles are projected against the surface to be coated forming adherent splats.

[0007] In the flame spray coating process, oxygen and a fuel such as acetylene are combusted in a torch. Powder, wire or rod is injected into the flame where it is melted and accelerated. Particle velocities may reach about 300 m/sec. The maximum temperature of the gas and ultimately the coating material is a function of the flow rate and composition of the gases used and the torch design. Again, the molten particles are projected against the surface to be coated forming adherent splats.

[0008] Thermal spray coating processes have been used for many years to deposit layered coatings. These coatings consist of discrete layers of different composition and properties. For example, the coating may be a simple duplex coating consisting of a layer of a metal alloy such as nickel-20 chromium (compositions herein are in weight percent unless otherwise noted) adjacent to the substrate with a layer of zirconia over it. In this case the undercoat of nickel-chromium may be used to enhance the mechanical or thermal shock resistance of the coating system or to protect the substrate from corrosion. An increase in mechanical or thermal shock resistance may be achieved by adding a third layer of coating consisting of a mixture of nickel-chromium and zirconia between the pure nickel-chromium and zirconia layers. Alternatively, perhaps even better thermal or mechanical shock resistance could be achieved by using two or more intermediate layers, each with an increasing amount of zirconia, thus approximating a continuously graded structure. Recently some graded coatings have been referred to as "functionally graded" coating systems.

[0009] Prior to the present invention the general means of creating graded coating structures was to intermittently change the powder, wire or rod coating material composition being fed to the thermal spray device or devices if more than one was used to deposit the various layers. The coating deposition parameters and in some cases the coating thermal spray device were changed with each layer to obtain the desired coating structure of the particular layer composition. In most cases this meant that the coating process had to be stopped; the powder, wire, or rod feeders and composition changed; the deposition parameters such as electrical power, gas flows, and gas compositions changed; the process restarted; and new coating qualified before the coating of the substrate could be continued. All of this added very substantial time and cost to the coating process. Moreover, the time between coating layers generally tends to decrease the bond strength between layers and the over-all strength of the coating system.

[0010] It is an object of the present invention to provide a novel process for the production of thermal spray coatings that are graded in composition and/or properties.

[0011] It is a further object of the present invention to provide novel coatings that are graded in composition and/or density properties produced by the said novel process.

[0012] It is also a further object of the present invention to provide articles with graded coatings produced using the novel process of the invention.

[0013] It is a further object of this invention to provide a process for the deposition of graded coatings with greater cohesive strength than can be achieved using the multiple layer coatings of the present art.

[0014] It is also a further object of this invention to provide graded coatings with greater cohesive strength than that of the graded coatings of the present art. The higher cohesive strength of the coatings of this invention is believed to be the result of the smoother transition in composition and properties achieved with the process of this invention and the minimal time between the deposition of the layers of the coating.

Summary of the Invention

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[0015] The invention relates to a process for producing a graded thermal spray coated layer on a substrate comprising the feeding of a mixture of at least two coating materials to a thermal spray device and varying at least one of the deposition parameters of the thermal spray device during the deposition operating thereby varying the composition of the deposited coating material to produce a graded coated heterogeneous layer on the substrate. The thermal spray device for the process of this invention has parameters that can control or monitor the temperature of the depositing coating material and the velocity of the coating material particles.

[0016] The invention also relates to the deposition by means of the novel coating process of this invention, of unique coating structures with smoothly varying gradations in composition and/or density properties. Since the changes in deposition parameters can be made while the coating is being continuously deposited, the gradation or changes in composition and/or density properties are very smooth. If the coating is being continuously deposited, the gradation or changes in composition and properties are very smooth. If the coating is deposited without moving the gun or torch

and the substrate is stationary as well, the coating gradation will be continuous; i.e., without discrete changes as a function of thickness. In most cases, however, the coating device and substrate can be moved relative to each other and the coating is deposited in multiple layers. Using the process of this invention, each layer may be slightly different than the preceding or succeeding layer. The time between layers is only dependent on the size of the substrate and the traverse rate (the relative rate of motion between the coating device and the substrate), since coating is being deposited continuously by the coating device. The difference between layers is a function of the rate of change in deposition parameters and the traverse rate. The smoothness of the gradation is then a function of the thickness of the individual layers which can be made very thin. The total thickness of the coating and of each zone is a function of the requirements of the application. The total thickness of the coating is typically in the range of 100 to 500 microns, but may be thicker or thinner if it is necessary to satisfy the specific requirements of the application.

[0017] This invention also relates to articles with the graded coatings of this invention. Such articles include those requiring coatings with graded properties to enhance the coating's mechanical, thermal, or electrical properties. As illustrative, but not limited, examples of these coated articles include (a) those that require a very hard, brittle surface such as an oxide or carbide graded to a tough, ductile metallic layer adjacent to the substrate to provide more impact resistance and higher bond strength of the coating to the substrate; (b) those that require a thermal barrier outer layer such as an oxide graded to a metallic layer adjacent to the substrate that provides oxidation resistance and thermal shock resistance; and (c) those that require a wear and corrosion resistant outer surface graded to an oxide layer adjacent to the substrate for electrical resistance.

[0018] As used herein, a graded coated layer shall mean at least one layer comprising a mixture of at least two coating materials in which at least one of the coating materials varies in composition to produce a heterogeneous coated layer. Also as used herein, composition of the coating material shall also include density of the coating material. [0019] Graded coatings, as defined herein, are used extensively to enhance the mechanical impact resistance, thermal shock resistance, and corrosion resistance of a coating system, as well as for other purposes. Occasionally graded coatings are used to allow thicker coatings to be deposited than would otherwise be possible. Most commonly, the layer of coating next to the substrate is a metallic alloy, and the outermost coating layer is an oxide or cermet. The metallic layer bonds better to the substrate and the oxide or cermet than the cermet or oxide directly to the substrate. It also may improve the mechanical impact resistance and other properties of the total coating by providing a layer of intermediate mechanical properties such as elastic modulus. Other factors such as stress relief through creep of the metallic layer may also be important. An example of this type of system is the use of a nickel based alloy under a tungsten carbide cobalt coating used to repair worn machine components. The thermal shock resistance of a coated system may also be increased with a metallic intermediate layer by increasing the bond strength of the system and providing an intermittent coefficient of thermal expansion between a metallic substrate and an oxide outer coating. This type of coating is frequently used in thermal barrier coating system where the metallic alloy undercoat is also used to protect the substrate from oxidation or other forms of corrosion. A typical example of a thermal barrier coating system, such as aircraft components such as aircraft blades, uses a cobalt-nickel-chromium-aluminum-yttrium metallic alloy undercoat and a zirconia-yttria outer coating. The porosity inherent in the metallic coating is sealed by heat treatment after deposition to provide a corrosion resistant barrier to protect the substrate, frequently a superalloy turbine blade or vane in a gas turbine engine.

[0020] In accordance with this invention the properties of many graded coating systems can be improved by increasing the number of layers of coatings with increasing amounts of the material of the final outer layer. This results in a smoother transition in properties as the number of layers increases.

[0021] The invention relates to a novel thermal spray process in which a mixture of coating materials is fed to a thermal spray gun or torch and the rate of deposition of the individual components of the mixture varied in a controlled manner by varying the thermal parameters of the thermal spray process gases. As a simple, but not limited example, consider a coating material that consists of two components, A and B, with different properties including melting point, size, shape, heat capacity, and thermal absorption characteristics. The thermal spray deposition parameters may be initially set to optimize the deposition rate or efficiency of A and not B and then changed gradually to optimize the deposition rate or efficiency of B. Thus the deposited coating would have a gradation in composition from predominantly A to A+B to predominately B. In addition to gradations in composition, gradations in other properties such as density can also be made by changing the deposition parameters. While this invention includes coatings that are continuously graded in composition and properties it also includes coatings in which one or more layers of the coating is maintained constant for a given thickness. Any and all of these variations in gradations will be encompassed in the term "graded" as used herein. The coating material is usually fed to the thermal spray device in the form of powder, although one or more of the constituents could be fed in the form of wire or rod. When two or more of the constituents of the coating material are in the form of powder they may be blended mechanically and fed from a single powder dispenser to the thermal spray device or fed individually or in partial blends from two or more powder dispensers to the thermal spray device. The coating material may be fed to the thermal spray device internally as in most detonation gun and high velocity oxy-fuel devices or externally as in many plasma spray devices. The changes in deposition parameters including gas composition and flow rates, power levels, and coating material injection rates may be changed during the deposition process either manually by the equipment operator or automatically by computer control.

Detailed Description

[0022] This invention comprises the process for depositing a graded coating using a thermal spray coating process in which a mixture of two or more powders is fed to the thermal spray device and the deposition parameters of the device are changed continuously or discontinuously. The result is that the deposition rate or efficiency of one or the powder constituents is initially dominant and, as the operating parameters are changed, the deposition rate of another constituent more or less gradually becomes dominant. An example, not intended to be limiting, is the deposition of a mixture of two materials with differing melting points such as a metallic alloy or an oxide. If a coating with a gradation from a metal layer next to the substrate to an oxide layer on the external surface of the coating is required, the operating parameters of the device would be initially set to provide a relatively low thermal content in the gas stream, sufficient to melt or nearly melt the metallic powder, but not the oxide. Under these conditions, the deposition rate of the metal would be very high while that of the oxide would be very low or nearly nil. The deposition parameters of the thermal spray device would then be changed to increase the thermal content of the gas stream so that more and more of the oxide would be heated to near or above its melting point. This would increase the deposition rate of the oxide and the coating composition would have a higher and higher fraction of oxide. As the thermal content of the gas stream increased to a sufficiently high level, the deposition rate of the metal would decline due to vaporization of the metal or due to the superheating of the particles to such a degree that they would splatter on impact rather than adhere to the surface being coated. The resulting coating may thus have a continuous gradation in composition from almost pure metal to almost pure oxide. If it were desired to have a constant composition for some portion of the thickness of the coating, the deposition parameters of the thermal spray device could be held constant for a period of time. Thus, continuing with the example of the metal-to-oxide coating, it would be possible to have a virtually pure metallic layer of some thickness adjacent to the substrate, followed by a graded zone, followed by a virtually pure oxide layer. This example could be used to produce a superior thermal barrier coating.

[0023] In those situations in which the thermal spray device is a detonation gun, the thermal content of the gas stream in the gun, as well as the velocity of the gas stream, can be varied by changing the composition of the gas mixtures. Both the fuel gas composition and the ratio of fuel to oxidant can be varied. The oxidant is usually oxygen. In the case of detonation gun deposition, the fuel is usually acetylene. In the case of Super D-Gun deposition, the fuel is usually a mixture of acetylene and another fuel such as propylene. The thermal content can be reduced by adding a neutral gas such as nitrogen.

[0024] In those situations in which the thermal spray device is a high velocity oxy-fuel torch or gun, the thermal content and velocity of the gas stream from the torch or gun can be varied by changing the composition of the fuel and the oxidant. The fuel may be a gas or liquid as described above. The oxidant is usually oxygen gas, but may be air or another oxidant.

[0025] In those situations in which the thermal spray device is a plasma spray torch, the thermal content of the plasma gas stream can be varied by changing the electrical power level, gas flow rates, or gas composition. As described above, argon is usually the base gas, but hydrogen, nitrogen, and helium are frequently added. Some plasma spray devices use liquid water rather than a gas to create the plasma stream. The velocity of the plasma gas stream can also be varied by changing the same parameters.

[0026] Variations in gas stream velocity from the thermal spray device can result in variations in particle velocities and hence dwell time of the particle in flight. This affects the time the particle can be heated and accelerated and, hence, its maximum temperature and velocity. Dwell time is also affected by the distance the particle travels between the torch or gun and the surface to be coated.

[0027] The specific deposition parameters used with any of the thermal spray devices depend on both the characteristics of the device and the materials being deposited. The rate of change or the length of time the parameters are held constant are a function of both the required coating composition gradation, the rate of traverse of the gun or torch relative to the surface being coated, and the size of the part. Thus a relatively slow rate of change when coating a large part may be the equivalent of a relatively large rate of change when coating a small part.

[0028] This invention also comprises graded coatings made by the process of this invention. These coatings have two or more constituents. The coatings of this invention include, but are not limited to, coatings that are virtually continuously graded from the composition of one constituent in the powder mixture to that of another constituent and also coatings graded from one constituent to another with zones of virtually constant composition for a given thickness. The gradation may have a very sharp transition zone such that the coating's composition changes from A to B with little discernable intermediate mixed zone. Alternatively, the transition may be very gradual from A to A+B to B with the A+B zone gradually changing from predominately A to predominantly B. More than two constituents may be included in the gradations. Thus, the gradation may be from A to B to C or may be from A to B+C, or from A to A+B to B+C, etc. In

addition to gradations in composition, gradations in properties such as density can be made by changing the deposition parameters, either simultaneously while changing composition or independently while holding the composition constant for a given thickness of the coating.

[0029] The coatings of this invention may consist of metallic, ceramic, cermet, or polymeric materials, alloys thereof, or any combination of these. The coatings of this invention may be used for a wide variety of purposes including, but not limited to, wear resistance, corrosion resistance, thermal characteristics, mechanical characteristics, optical characteristics, or electrical characteristics or for any combination of these. A few examples, not intended to be limited in any way, include their use in the following ways: (a) as wear resistant surfaces where the gradation in properties may be from a metallic zone adjacent to a metallic substrate to hard wear resistant cermet composition on the exterior surface with the metallic zone and gradation serving to increase bond strength and ameliorate the mechanical stresses imposed in the coating in service; (b) as thermal barriers where the coating is graded from a zone of metallic coatings adjacent to the metallic substrate to an oxide with low thermal conductivity where the metallic layer provides corrosion protection and better bond strength and the gradation provides better thermal shock resistance; or (c) as electrically insulated and wear resistant coatings where the composition of the coating adjacent to a metallic substrate is an electrical insulator and the gradation is to a more wear resistant cermet surface where the gradation provides better bond strength and ameliorates thermal stresses placed on the coating in service.

[0030] The coated articles of this invention comprise those articles having a graded coating produced using the process of this invention. Said articles may include a very wide variety of components of manufacturing equipment or end products of production. A few examples intended to illustrate such articles, but not to be limiting in any way, include the following: (a) components requiring wear resistance in service that includes high mechanical loading where the coating may include a gradation from a relative tough, but ductile, metallic zone adjacent to a metallic substrate to a wear resistant cermet exterior zone with the gradation providing better bond strength and dissipation of the mechanical stresses placed on the coating's surface; (b) components requiring a thermal barrier to protect them from excessive heat or thermal shock or to increase the thermal efficiency of a process where the coating is graded from a metallic to an oxide as noticed above; or (c) components requiring electrical insulation with either the insulating layer being adjacent to the substrate with gradation to a more wear resistant surface as noted above or where the coating is graded from a metallic adjacent to substrate with gradation to an insulating oxide on the exterior surface with the gradation providing better bond strength and amelioration of imposed thermal or mechanical stresses.

EXAMPLE ...

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[0031] A wear and corrosion resistant coating consisting of chromium carbide plus a nickel-chromium alloy known as nichrome that was electrically isolated from a metallic substrate that would also withstand thermal shock was required. An intermediate layer of alumina was chosen for the electrical isolation. Repeated attempts to apply the two materials as individual layers according to the prior art via detonation gun deposition were unsuccessful. The chromium carbide plus nichrome coating either would not adhere properly or would spall during testing. The introduction of an intermediate layer of a mixture of alumina plus chromium carbide/nichrome to form a graded coating according to the prior art was similarly unsuccessful. Surprisingly, a coating that consisted of a first zone adjacent to the substrate of virtually pure alumina, a second zone of alumina plus chromium carbide plus nichrome, and a third zone of chromium carbide plus nichrome deposited using the process of the present invention was very successful. A detonation gun made by Praxair Surface Technologies, Inc. was used to produce the graded coating of this invention. The powder mixture used consisted of 50 volume percent pure alumina blended with a previously mechanically blended mixture of 80 weight percent Cr₃C₂ plus 20 weight percent of nichrome (an alloy of 80 weight percent nickel plus 20 weight percent chromium). The particle sizes of the constituents were:

Alumina - 5 to 44 microns
Chromium carbide - 5 to 44 microns, and
Nichrome - 5 to 53 microns

[0032] The deposition parameters for each of the zones were:

	First Zone	Second Zone	Third Zone
Acetylene	4.60	3,59	2.20
Oxygen	6.50	4.66	2.53
Nitrogen	0.00	2.75	6.27
Total gas flow, cfm	11.10	11.00	11.00

(continued)

	First Zone	Second Zone	Third Zone
Oxygen/carbon ratio	1.41	1.30	1.15
Nitrogen dilution, %	0.00	25.00	57.00

[0033] The thickness of the zones of the coating were, in microns:

	First Zone	Second Zone	Third Zone
į	73.5	112.5	122.5
į	to 85.8	to 147	to 147

[0034] The high oxygen to carbon ratio and no dilution result in a very high detonation temperature and thermal content in the gas stream for the first zone. Thus a very high deposition rate or efficiency for the alumina constituent and very low or nil deposition rate or efficiency for the other constituents were achieved. The intermediate oxygen to carbon ratio and 25% nitrogen dilution resulted in an intermediate gas temperature and heat content for the second zone yield. Thus approximately equal rates of deposition or affiances for the alumina and the combination of chromium carbide plus chrome were achieved. The relatively low oxygen to carbon ratio and high (57%) nitrogen dilution resulted in a low gas stream temperature and low heat content. Thus a low deposition rate or efficiency for the alumina was achieved and a predominantly chromium carbide plus nichrome zone was created. This coating exhibited very high coherent strength and withstood testing of the coating in thermal shock.

[0035] Those skilled in the art can readily find many and widely varying uses for the unique and surprising useful processes, coatings, and coated articles of this invention. As many possible embodiments of this invention may be made without departing from the scope of the invention, it is understood that all matter set forth herein is to be interpreted as illustrative and in no way limiting.

Claims

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- A process for producing a graded thermal spray coated layer on a substrate comprising the feeding of a mixture
 of at least two coating materials to a thermal spray device and varying at least one of the deposition parameters
 of the thermal spray device during the depositor operation thereby varying the composition of the deposited coating
 material to produce a graded coated layer on the substrate.
- 2. The process of claim 1 whereby the parameters of the thermal spray device to be varied are selected from the group consisting of varying the temperature of the depositing coating material and varying the velocity of the coating material as it contacts the substrate.
- The process of claim 2 wherein the coating material is heated to about its melting point to form droplets of the coating material and accelerating the droplets in a gas flow stream to contact the substrate.
- 4. The process of claim 3 wherein the temperature parameters of the coating material can be varied with at least one function selected from the group consisting of the temperature and enthalpy of the gas stream; the composition and thermal properties of the coating droplets; the size and shape distributions of the droplets; the mass flow rate of the droplets relative to the gas flow rate; and the time of transit of the droplets to the substrate.
- 5. The process of claim 3 wherein the velocity parameters of the coating material can be varied with at least one function selected from the group consisting of gas flow rate; size and shape distribution of the droplets; and mass injection rate and density of the droplets.
 - 6. The process of claim 1 wherein the materials in the mixture fed to the thermal spray device are selected from the group comprising metallic, ceramic, cermet, polymeric materials and alloys thereof.
- 7. A graded coated substrate having at least one graded coated layer and said graded coated layer is formed by producing a graded thermal spray coated layer on a substrate comprising the feeding of a mixture of at least two coating materials to a thermal spray device and varying at least one of the deposition parameters of the thermal

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spray device during the depositor operation thereby varying the composition of the deposited coating material to produce a graded coated layer on the substrate.

- 8. The graded coating substrate of claim 7 wherein the graded coated layer is made from material selected from the group comprising metallic, ceramic, cermet, or polymeric materials, or alloys thereof.
 - 9. The graded coated substrate of claim 7 wherein the graded coated layer is comprised of alumina and chromium carbide with nichrome.
- 10. The graded coated substrate of claim 7 is a blade.

(11) EP 1 160 348 A3

(12)

EUROPEAN PATENT APPLICATION

(88) Date of publication A3: 29.10.2003 Bulletin 2003/44

(51) Int Cl.7: C23C 4/02

- (43) Date of publication A2: 05.12.2001 Bulletin 2001/49
- (21) Application number: 01112406.2
- (22) Date of filing: 21.05.2001
- (84) Designated Contracting States:

 AT BE CH CY DE DK ES FI FR GB GR IE IT LI LU

 MC NL PT SE TR

 Designated Extension States:

 AL LT LV MK RO SI
- (30) Priority: 22.05.2000 US 576439
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- (54) Process for producing graded coated articles
- (57) The invention is directed to a process for improving the separation of sodium bicarbonate brine from crude tall oil or of sulphate brine from soap oil in the recovery process of a pulp mill. In the neutralisation/

acidulation of crude tall oil soap (6) or soap oil (9) with carbon dioxide (5) respectively sulphuric acid (11) a water solution (15) having an increased density is used.



EUROPEAN SEARCH REPORT

Application Number

EP 01 11 2406

		ERED TO BE RELEVANT	T	
Category	Citation of document with in of relevant pass	ndication, where appropriate, ages	Relevant to claim	CLASSIFICATION OF THE APPLICATION (INLCI.7)
X	US 4 588 607 A (ALF 13 May 1986 (1986-0 + claims 1,3-6,10 +		1-8,10	C23C4/02
X	GB 2 130 250 A (GLY 31 May 1984 (1984-0 * page 2, line 60 -		1,3,6-8	
X	DE 37 21 008 A (GLY & LOOS) 20 October * column 10, line 5 1,7,10,15-17,19,23	- line 28; claims	1,3,6-8	
X	US 3 912 235 A (KEN 14 October 1975 (19 + abstract; claim 1	75-10-14)	1,2,6-8	
X	US 5 988 488 A (KEV 23 November 1999 (1 + claims 1,2 *		1,3,6-8	
X	US 4 751 099 A (MAS 14 June 1988 (1988- + claims 1-5; examp	06-14)	1,6-8,1	O SEARCHED (INI.CI.7) C23C
X	EP 0 816 527 A (FOR 7 January 1998 (199 + claim 1 +	D MOTOR COMPANY) 8-01-07)	1,3-8	
X	US 3 413 136 A (ROG 26 November 1968 (1 * column 2, line 67 claim 1 *	ER C. EMANUELSON) 968-11-26) - column 3, line 21;	1-8,10	
X	EP 0 217 991 A (REP 15 April 1987 (1987 * claims 1,11,13,14	-04-15) * 	1,3,6-8	
				40 VO 10 VO
	The present search report has	been drawn up for all claims		
	Place of swarch	Date of completion of the search		Examiner
	THE HAGUE	9 September 2003	E1	sen, D
X:pax Y:pax doc A:ted O:not	CATEGORY OF CITED DOCUMENTS tlouterly relevant if balish alone tlouterly relevant if combined with anot unmore of the aarne category hnological background 	L: document about	ocument, but put ate in the application for other reason	blished on, or on on



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Application Number

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alegory	Citation of document with income of relevant passa			evant laim	CLASSIFICATION OF THE APPLICATION (Int.CL7)
x	US 5 814 152 A (STEP 29 September 1998 (1 * column 11, line 15 * abstract *	HEN L. THALER) 998-09-29)		-8,10	AFERMION (MCCLT)
X	US 5 612 099 A (STEP 18 March 1997 (1997- * column 11, line 6 * abstract *	03-18)	1,3-	-8,10	
X	EP 0 290 368 A (UNIT CORPORATION) 9 Novem * column 6, line 42 * abstract; claims 1	ber 1988 (1988-11-09 - line 51 *	1,6	-8	
X	PATENT ABSTRACTS OF vol. 1998, no. 02, 30 January 1998 (199 & JP 09 263927 A (107 October 1997 (1997 * abstract *	98-01-30) BYOTA MOTOR CORP),	1,6	- 8 ;	TECHNICAL FIELDS SEARCHED (Int.Cl.7)
X	WO 93 05194 A (TECHN 18 March 1993 (1993- + claims 1,19,21-23	-03-18)	1-3	,5-8	SENSIES (Massir)
X	US 4 613 259 A (LOU) 23 September 1986 (1 * abstract *		1,6	-8	
A	DE 34 35 748 A (SIEN 10 April 1986 (1986- + claims 1-4; figure	-04-10)	2-6		
	The present search report has b	een drawn up for all claims			
			1	Elsi	en, D
X:per Y:par doc	CATEGORY OF CITED DOCUMENTS ricularly relevant if taken alone ricularly relevant if combined with anoth zument of the same category fundosjical background	L : document o	rinciple under int document ing date cited in the ap	tying the in but public plication seasons	mention shed on, or



EUROPEAN SEARCH REPORT

Application Number

EP 01 11 2406

ategory	Citation of document with in of relevant pass	dication, where appropriate,	Relevant	CLASSIFICATION OF T	HE
1	PATENT ABSTRACTS OF vol. 016, no. 090 (5 March 1992 (1992-	JAPAN M-1218), 03-05) 0CALO CO LTD;OTHERS:	to claim	APPLICATION (InLCI.	<i>(</i>)
			•		
	·	·			
	·			TECHNICAL FIELDS	
				SEARCHED (Int.C	:1.7)
:		·			
			•		
		-			

	The present search report has	been drawn up for all claims			
	Place of search	Date of completion of the seeich	T	Examiner	
	THE HAGUE	9 September 2003	Els	en, D	
X:par Y:par doc A:ted	CATEGORY OF CITED DOCUMENTS ticularly relevant if taken alone ticularly relevant if combined with a not tament of the same category inhological background n-written disclosure	E : eafter patent docum after the flang date O : document clied in th L : document cited for o	nderlying the l nent, but publi ne epplication of epplications	invention shed on, or	

4

ANNEX TO THE EUROPEAN SEARCH REPORT ON EUROPEAN PATENT APPLICATION NO.

EP 01 11 2406 ---

This annex lists the patent family members relating to the patent documents cited in the above-mentioned European search report. The members are as contained in the European Patent Office EDP file on The European Patent Office is in no way liable for these particulars which are merely given for the purpose of information.

09-09-2003

	Patent documer cited in search rep		Publication date		Patent fam member(s		Publication date
us	4588607	A	13-05-1986	DE	3564453	D1	22-09-1988
••	100007	••	10 00 1700	EP	0183638		04-06-1986
				ĴΡ	1776831	–	28-07-1993
				JP	4048867	-	07-08-1992
				JP	61143576	_	01-07-1986
					01143570		01-07-1980
GB	2130250	Α	31-05-1984	DE	3242543	- · ·	24-05-1984
				BR	8305987	- •	10-07-1984
				FR	2536309	–	25-05-1984
				JP	59096945	Α	04-06-1984
DE	3721008	Α	20-10-1988	DE	3721008	Al	20-10-1988
				DE	8717379	U1	20-10-1988
US	3912235	A	14-10-1975	NONE	********		
US	5988488	A	23-11-1999	NONE	*		
US.	4751099	Α	14-06-1988	JP	1830726	C	15-03-1994
•	1,21033	**	14 00 1300	JP	5036229	-	28-05-1993
				JP	62156938		11-07-1987
				DE	3679975	• •	01-08-1991
				EP	0229522		22-07-1987
FP.	816527	Α	07-01-1998	US	5958521	Δ	28-09-1999
	010027	••	0, 41 1550	CA	2208398		21-12-1997
				DE	69702576		24-08-2000
					69702576		12-04-2001
				EP	0816527		07-01-1998
US	3413136	A	26-11-1968	US	3545944	A	08-12-1970
EP	217991	A	15-04-1987	EP	0217991	A1	15-04-1987
				AU	4054485		03-10-1985
				JP	62093359	A	28-04-1987
US	5814152	Α	29-09-1998	us	5612099	A	18-03-1997
				MO	9832891		30-07-1998
US	5612099	Α	18-03-1997	WO.	9832891	Δ1	30-07-1998
-	0016077	•	20 00 1937	US	5814152		29-09-1998
				AU	2112797		18-08-1998
	200260	4	00 11 1000		4700041		00.11.1011
t٢	290368	A	09-11-1988	US	4783341		08-11-1988
				DE DE	3873972		01-10-1992
				DE	3873972	12	14-01-1993

For more details about this annex : see Official Journal of the European Patent Office, No. 12/82

ANNEX TO THE EUROPEAN SEARCH REPORT ON EUROPEAN PATENT APPLICATION NO.

EP 01 11 2406

This annex lists the patent family members relating to the patent documents cited in the above-mentioned European search report. The members are as contained in the European Patent Office EDP file on The European Patent Office is in no way liable for these particulars which are merely given for the purpose of information.

09-09-2003

Patent documer cited in search rep		Publication date		Patent family member(s)	Publication date
EP 290368	A		EP JP JP JP	0290368 A2 1631445 C 2055501 B 63293152 A	09-11-1988 26-12-1991 27-11-1990 30-11-1988
JP 09263927	Α	07-10-1997	NONE		
WO 9305194	Α	18-03-1993	.WO US	9305194 A1 5362523 A	18-03-1993 08-11-1994
US 4613259	A	23-09-1986	DE EP JP JP JP	3565944 D1 0185604 A1 1728093 C 4009106 B 61141951 A	08-12-1988 25-06-1986 19-01-1993 19-02-1992 28-06-1986
DE 3435748	A	10-04-1986	DE	3435748 A1	10-04-1986
JP 03272959	A	04-12-1991	NONE		
	•				

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